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## **INFORMATION DISCLOSURE STATEMENT BY APPLICANT**

(Use as many sheets as necessary)

Sheet 1 of 1

<i>Complete if Known</i>	
Application Number	10/573,890
Filing Date	03-29-06
First Named Inventor	Kazutaka Nakamoto
Art Unit	1625
Examiner Name	Patricia L. Morris
Attorney Docket Number	3939-01118PUS1

**U.S. PATENT DOCUMENTS**

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**FOREIGN PATENT DOCUMENTS**

FOREIGN PATENT DOCUMENTS						
Examiner Initial *	Cite No. 1	Foreign Patent Document		Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Country <sup>3</sup>	Number <sup>4</sup> Kind Code (if known) <sup>5</sup> Code			
	12	EP	0 124 154	11-07-1984		
	13	WO	03-091226	11-06-2003		
	14	WO	03-091227	11-06-2003		
	15	JP	2001-527083	12-25-2001		
	16	WO	2004-014366	02-19-2004		
	17	JP	2005-526751	09-08-2005		
	18	JP	2006-519247	08-24-2006		
	19	WO	2009-084621	07-09-2009		
	20	JP	52-94935	11-09-1993		
	21	JP	59-073575	04-25-1984		

Examiner Signature	Date Considered	
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3. Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3.4). For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. 5. Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST.16 if possible. 6. Applicant is to place a check mark here if English language Translation is attached.

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<sup>10</sup> See, for example, the discussion of the 1993 Constitutional Conference in the *Journal of African Law* (1994), 38(1), 1–20.

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## INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 2 of 3

**Complete if Known**

<b>Application Number</b>	10/573,890
<b>Filing Date</b>	03-29-06
<b>First Named Inventor</b>	Kazutaka Nakamoto
<b>Art Unit</b>	1625
<b>Examiner Name</b>	Patricia L. Morris
<b>Attorney Docket Number</b>	3939-0118PUS1

## U.S. PATENT DOCUMENTS

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**FOREIGN PATENT DOCUMENTS**

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Examiner Signature	Date Considered	
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3. Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3.) 4. For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. 5. Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST.16 if possible. 6. Applicant is to place a check mark here if English language Translation is attached.

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INFORMATION DISCLOSURE  
STATEMENT BY APPLICANT

(Use as many sheets as necessary)

Sheet

3

of

3

Complete if Known	
Application Number	10/573,890
Filing Date	03-29-06
First Named Inventor	Kazutaka Nakamoto
Group Art Unit	
Examiner Name	

Attorney Docket Number 3939-0118PUS1

## NON PATENT LITERATURE DOCUMENTS

Examiner initial *	Cite No. 1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T 2
23	CHANG et al., "Synthesis and Structure-Activity Relationships of Quaternary Ammonium Cephalosporins with 3-Pyrazolylpyridinium Derivatives," <i>Bioorganic &amp; Medicinal Chemistry Letters</i> (2000) Vol. 10, No. 11, pp. 1211-1214		
24	CONNORS et al., "Prodrugs in medicine," Overview, <i>Biologicals &amp; Immunologicals, Exp. Opin. Ther. Patents</i> , Vol. 5, No. 9, 1995, pp. 873-885		
25	Copy of an Office Action from co-pending U.S. Patent Application No. 11/589,128, mailed May 7, 2009		
26	HATA, "New Approaches to Antifungal Drugs for the Treatment of Fungal and Protozoal Infections, Rauconazole and Beyond: New Targets and Pre-clinical Strategies," The SMI's 12th Annual Conference, Superbugs and Superdrugs, March 18, 2010, Crowne Plaza London - St. James, 44 pages		
27	International Search Report dated May 20, 2008 for corresponding International Application No. PCT/JP2008/057851		
28	ISHIKAWA et al., "TAK-599, a Novel N-Phosphon Type Prodrug of Anti-MRSA Cephalosporin T-91825: Synthesis, Physicochemical and Pharmacological Properties," <i>Bioorganic &amp; Medicinal Chemistry</i> , Vol. 11, pp. 2427-2437, (2003)		
29	LUKEVICS et al., "Synthesis and cytotoxicity of silyl- and carbonyl-substituted isoxazoles," <i>Chemistry of Heterocyclic Compounds</i> (2000) Vol. 36, No. 10, pp. 1226-1231		
30	PLATE et al., "Synthesis and Muscarinic Activities of 3-(Pyrazolyl)-1,2,5,6-tetrahydropyridine Derivatives," <i>Bioorganic &amp; Medicinal Chemistry Letters</i> (1996) Vol. 4, No. 2, pp. 227-237		
31	Supplementary European Search Report dated February 6, 2009 for corresponding European Application No. 04788159.4		
	TANAKA et al., "An Effective Lewis Acid-Mediated 1,3-Dipolar Cycloaddition of Nitrile Oxide Using Acetylene: Synthesis of a (2-Aminopyridin-3-yl)isoxazole Derivative and Its Application to Novel Antifungal Agents," pp. 1-8		
32	VRZHESCHCH et al., "Supercooperativity in platelet aggregation: Substituted pyridyl isoxazoles, a new class of supercooperative platelet aggregation inhibitors," <i>FEBS Letters</i> (1994) Vol. 351, No. 2, pp. 168-170		

Examiner Signature	/Patricia L. Morris/ (07/29/2010)	Date Considered

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